STN Columbus

* * *	* *	* *	* *	* Welcome to STN International * * * * * * * * *	
NEWS	1			Web Page for STN Seminar Schedule - N. America	
NEWS	2	NOV	21	CAS patent coverage to include exemplified prophetic	
112110	-	1101		substances identified in English-, French-, German-,	
				and Japanese-language basic patents from 2004-present	
NEWS	3	NOV	26	MARPAT enhanced with FSORT command	
NEWS	4	NOA		CHEMSAFE now available on STN Easy	
NEWS	5	NOV	26	Two new SET commands increase convenience of STN	
				searching	
NEWS	6	DEC	01	ChemPort single article sales feature unavailable	
NEWS	7	DEC	12	GBFULL now offers single source for full-text	
				coverage of complete UK patent families	
NEWS	8	DEC		Fifty-one pharmaceutical ingredients added to PS	
NEWS	9	JAN	06	The retention policy for unread STNmail messages	
				will change in 2009 for STN-Columbus and STN-Tokyo	
NEWS	10	JAN	0 /	WPIDS, WPINDEX, and WPIX enhanced Japanese Patent	
			0.0	Classification Data	
NEWS	11	FEB	02	Simultaneous left and right truncation (SLART) added	
NEWS	1.2	FEB	0.2	for CERAB, COMPUAB, ELCOM, and SOLIDSTATE GENBANK enhanced with SET PLURALS and SET SPELLING	
NEWS		FEB		Patent sequence location (PSL) data added to USGENE	
NEWS		FEB		COMPENDEX reloaded and enhanced	
NEWS		FEB		WTEXTILES reloaded and enhanced	
NEWS		FEB		New patent-examiner citations in 300,000 CA/CAplus	
				patent records provide insights into related prior	
				art	
NEWS	17	FEB	19	Increase the precision of your patent queries use	
				terms from the IPC Thesaurus, Version 2009.01	
NEWS	18	FEB	23	Several formats for image display and print options	
				discontinued in USPATFULL and USPAT2	
NEWS	19	FEB	23	MEDLINE now offers more precise author group fields	
				and 2009 MeSH terms	
NEWS	20	FEB	23	TOXCENTER updates mirror those of MEDLINE - more	
115110	0.1		0.0	precise author group fields and 2009 MeSH terms	
NEWS	21	FEB	23	Three million new patent records blast AEROSPACE into	
NEWS	22	FEB	2.5	STN patent clusters USGENE enhanced with patent family and legal status	
MEMO	22	FED	23	display data from INPADOCDB	
NEWS	23	MAR	06	INPADOCDB and INPAFAMDB enhanced with new display	
112110	2.0		00	formats	
NEWS	2.4	MAR	11	EPFULL backfile enhanced with additional full-text	
				applications and grants	
NEWS	25	MAR	11	ESBIOBASE reloaded and enhanced	
NEWS	EXP	RESS		E 27 08 CURRENT WINDOWS VERSION IS V8.3,	
			AND	CURRENT DISCOVER FILE IS DATED 23 JUNE 2008.	
NEWS				N Operating Hours Plus Help Desk Availability	
NEWS				lcome Banner and News Items	
NEWS	IPC	3	ro:	r general information regarding STN implementation of IPC 8	
Enter NEWS followed by the item number or name to see nove on that					
Enter NEWS followed by the item number or name to see news on that specific topic.					
opect:		JUPI	-•		
A11	All use of STN is subject to the provisions of the STN Customer				
				ase note that this agreement limits use to scientific	
				for software development or design or implementation	
	of commercial gateways or other similar uses is prohibited and may				

of commercial gateways or other similar uses is prohibited and may result in loss of user privileges and other penalties.

FILE 'HOME' ENTERED AT 22:22:24 ON 13 MAR 2009

FILE 'USPATFULL' ENTERED AT 22:26:23 ON 13 MAR 2009
CA INDEXING COPYRIGHT (C) 2009 AMBRICAN CHEMICAL SOCIETY (ACS)

FILE 'USPATOLD' ENTERED AT 22:26:23 ON 13 MAR 2009
CA INDEXING COPYRIGHT (C) 2009 AMERICAN CHEMICAL SOCIETY (ACS)

ETT B. HODATAL PURENER AT 22-26-22 ON 12 MAR 2000

FILE 'USPAT2' ENTERED AT 22:26:23 ON 13 MAR 2009 CA INDEXING COPYRIGHT (C) 2009 AMERICAN CHEMICAL SOCIETY (ACS)

=> s tamsulosin L1 1288 TAMSULOSIN

LI 1200 IAMOULUSI

=> s tamsulosin/clm L2 226 TAMSULOSIN/CLM

=> s polymer? L3 1224605 POLYMER?

=> s polymer?/clm

L4 432182 POLYMER?/CLM

-> s (calcium or potassium or sodium or magnesium)

L5 1287207 (CALCIUM OR POTASSIUM OR SODIUM OR MAGNESIUM)

=> s (calcium or potassium or sodium or magnesium)/clm L6 305082 (CALCIUM OR POTASSIUM OR SODIUM OR MAGNESIUM)/CLM

-> s (multilayer or layer) L7 2063906 (MULTILAYER OR LAYER)

=> s (multilayer or layer)/clm L8 810896 (MULTILAYER OR LAYER)/CLM

=> s capsule?

L9 243183 CAPSULE?

=> s capsule?/clm

L10 27197 CAPSULE?/CLM

897 L1 AND L3

=> s 11 and 14 L11 242 L1 AND L4

=> s 11 and 13

=> s 15 and 112

L13 888 L5 AND L12

=> s 17 and 113 L14 600 L7 AND L13

=> s 19 and 114 L15 493 L9 AND L14

=> s 12 and 14 L16 50 L2 AND L4

=> s 16 and 116 L17 33 L6 AND L16

=> s 18 and 117 L18 6 L8 AND L17

=> s 110 and 118 L19 2 L10 AND L18

=> d 1-2

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L19 ANSWER 1 OF 2 USPATFULL on STN
Full Text
       2009:4272 USPATFULL
ΔN
       Controlled release tamsulosin hydrochloride formulation
IN
       Cheng, Xiu Xiu, Weston, FL, UNITED STATES
       Cheng, Xiufang, Weston, FL, UNITED STATES
       Watson Pharmaceuticals, Inc. (U.S. corporation)
US 20090004284 Al 20090101
PA
PT
       IIS 2007-821990
                             A1 20070626 (11)
AΤ
       Utility
       APPLICATION
FS
LN.CNT 567
       INCLM: 424/497.000
INCL
       INCLS: 424/490.000; 514/603.000
       NCLM: 424/497.000
NCL
       NCLS: 424/490.000; 514/603.000
       IPCI
               A61K0009-14 [I,A]; A61K0031-18 [I,A]
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L19 ANSWER 2 OF 2 USPATFULL on STN
Full Text
       2008:253184 USPATFULL
AN
ΤI
       Advanced drug development and manufacturing
IN
       Birnbaum, Eva R., Los Alamos, NM, UNITED STATES
       Koppisch, Andrew T., Flagstaff, AZ, UNITED STATES
       Baldwin, Sharon M., Santa Fe, NM, UNITED STATES
       Warner, Benjamin P., Los Alamos, NM, UNITED STATES McCleskey, T. Mark, Los Alamos, NM, UNITED STATES
       Stewart, Jeffrey Joseph, Los Alamos, NM, UNITED STATES
       Berger, Jennifer A., Los Alamos, NM, UNITED STATES
       Harris, Michael N., Los Alamos, NM, UNITED STATES
       Burrell, Anthony K., Los Alamos, NM, UNITED STATES
ΡI
       US 20080220441
                           A1 20080911
A1 20071010 (11)
ΑI
       US 2007-974156
       Continuation-in-part of Ser. No. US 2001-859701, filed on 16 May 2001, 
PENDING Continuation-in-part of Ser. No. US 2002-206524, filed on 25 Jul 2002, ABANDONBE Continuation-in-part of Ser. No. US 2003-621825, filed
RLI
       on 16 Jul 2003, Pat. No. US 6858148
PRAT
       IIS 2006-850594P
                            20061010 (60)
       Utility
DT
FS
       APPLICATION
LN.CNT 10199
INCL
       INCLM: 435/071.000
       INCLS: 436/501.000; 436/172.000; 436/086.000; 378/045.000
NCL
       NCLM: 435/007.100
       NCLS:
              378/045.000; 436/086.000; 436/172.000; 436/501.000
       IPCI
               G01N0033-53 [I,A]; G01N0021-76 [I,A]; G01N0033-68 [I,A];
               G01N0023-223 [I,A]; G01N0023-22 [I,C*]
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
=> d 118 1-6
L18 ANSWER 1 OF 6 USPATFULL on STN
Full Text
       2009:4272 USPATFULL
AN
ΤI
       Controlled release tamsulosin hydrochloride formulation
TN
       Cheng, Xiu Xiu, Weston, FL, UNITED STATES
       Cheng, Xiufang, Weston, FL, UNITED STATES
       Watson Pharmaceuticals, Inc. (U.S. corporation)
PA
ΡI
       US 20090004284 A1 20090101
AΙ
       US 2007-821990
                             A1 20070626 (11)
       Utility
       APPLICATION
LN.CNT 567
INCL
       INCLM: 424/497.000
       INCLS: 424/490.000; 514/603.000
       NCLM: 424/497.000
NCL
       NCLS: 424/490.000; 514/603.000
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CAS INDEXING IS AVAILABLE FOR THIS PATENT.
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L18 ANSWER 2 OF 6 USPATFULL on STN
Full Text
ΔN
       2008:354338 USPATFULL
       Solid form
IN
       Darmuzey, Olivia, Brussels, BELGIUM
       MacLeod, Graeme, Wezembeek Oppem, BELGIUM
Cengic, Dzenana, Brussels, BELGIUM
       US 20080311162
                         A1 20081218
PT
                            A1 20070516 (11)
AΤ
       US 2007-803825
DT
       Utility
       APPLICATION
FS
LN.CNT 1512
       INCLM: 424/401.000
INCL
       INCLS: 424/490.000; 514/263.340
       NCLM: 424/401.000
NCL
       NCLS:
              424/490.000; 514/263.340
       IPCI
              A61K0008-02 [I,A]; A61K0009-14 [I,A]; A61K0031-522 [I,A];
               A61K0031-519 [I,C*]; C11D0017-06 [I,A]
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L18 ANSWER 3 OF 6 USPATFULL on STN
AN
       2008:268489 USPATFULL
ΤI
       UROLOGICAL MEDICAL DEVICES FOR RELEASE OF THERAPEUTIC AGENTS
       Cheng, Eric, Bloomington, IN, UNITED STATES
IN
       Li, Jianmin, Lexington, MA, UNITED STATES
       Bucay-Couto, Weena, Burlington, MA, UNITED STATES
       Sanders, Scott, Hinsdale, IL, UNITED STATES
       Schuermann, James F., Natick, MA, UNITED STATES
       Sheu, Min-Shyan, Chelmsford, MA, UNITED STATES
PA
       Boston Scientific Scimed, Inc., Maple Grove, MN, UNITED STATES (U.S.
       corporation)
PΙ
       US 20080234659
                            A1 20080925
       US 2008-52037
                            A1 20080320 (12)
AΙ
PRAI
       US 2007-919081P
                            20070320 (60)
DT
       Utility
       APPLICATION
FS
LN.CNT 1180
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       INCLM: 604/523.000
       INCLS: 623/023.660
       NCLM: 604/523.000
NCL
       NCLS: 623/023.660
       IPCI
              A61M0025-00 [I,A]; A61F0002-04 [I,A]
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L18 ANSWER 4 OF 6 USPATFULL on STN
Full Text
AN
       2008:253184 USPATFULL
       Advanced drug development and manufacturing
       Birnbaum, Eva R., Los Alamos, NM, UNITED STATES
Koppisch, Andrew T., Flagstaff, AZ, UNITED STATES
IN
       Baldwin, Sharon M., Santa Fe, NM, UNITED STATES
       Warner, Benjamin P., Los Alamos, NM, UNITED STATES
McCleskey, T. Mark, Los Alamos, NM, UNITED STATES
       Stewart, Jeffrey Joseph, Los Alamos, NM, UNITED STATES
       Berger, Jennifer A., Los Alamos, NM, UNITED STATES
       Harris, Michael N., Los Alamos, NM, UNITED STATES
       Burrell, Anthony K., Los Alamos, NM, UNITED STATES
ΡI
       US 20080220441
                          A1 20080911
A1 20071010 (11)
ΑI
       US 2007-974156
RLI
       Continuation-in-part of Ser. No. US 2001-859701, filed on 16 May 2001,
       PENDING Continuation-in-part of Ser. No. US 2002-206524, filed on 25 Jul
       2002, ABANDONED Continuation-in-part of Ser. No. US 2003-621825, filed
       on 16 Jul 2003, Pat. No. US 6858148
       US 2006-850594P
                           20061010 (60)
PRAT
DT
       Utility
FS
       APPLICATION
LN.CNT 10199
TNCI.
       INCLM: 435/071.000
       INCLS: 436/501.000; 436/172.000; 436/086.000; 378/045.000
NCT.
       NCLM: 435/007.100
       NCLS: 378/045.000; 436/086.000; 436/172.000; 436/501.000
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IC
       IPCI
              G01N0033-53 [I.A]; G01N0021-76 [I.A]; G01N0033-68 [I.A];
              G01N0023-223 [I,A]; G01N0023-22 [I,C*]
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L18 ANSWER 5 OF 6 USPATFULL on STN
   Text
AN
       2008:86578 USPATFULL
TT
       Tamsulosin controlled-release tablet
TN
       Gan, Yong, Huairou, CHINA
       Zhou, Xinteng, Huairou, CHINA
PA
       Ocean Star International, Inc., Snowville, UT, UNITED STATES (non-U.S.
       corporation)
       US 20080075775
US 2006-580215
                           A1 20080327
A1 20061011 (11)
AΙ
       CN 2006-10153091
PRAI
                           20060922
       Utility
DT
FS
       APPLICATION
LN.CNT 720
INCL
       INCLM: 424/473.000
NCL
       NCLM: 424/473.000
             A61K0009-24 [I,A]
A61K0009-24 [I,C]; A61K0009-24 [I,A]
       IPCR
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L18 ANSWER 6 OF 6 USPATFULL on STN
       2007:140489 USPATFULL
AN
TT
       Sustained release formulations
IN
       Cho, Seong Hwan, Suwon-city, KOREA, REPUBLIC OF
       Ku, Jeong, Yongin-city, KOREA, REPUBLIC OF
       Lim, Dong Kwon, Yongin-city, KOREA, REPUBLIC OF
       Cheon, Jun Hee, Suwon-city, KOREA, REPUBLIC OF
       An, Tae Kun, Yongin-city, KOREA, REPUBLIC OF
       Ko, Jae Kyoung, Incheon-city, KOREA, REPUBLIC OF
       Youn, Yong Sik, Yongin-city, KOREA, REPUBLIC OF
       Park, Choong Sil, Icheon-city, KOREA, REPUBLIC OF
Suh, Hea Ran, Icheon-city, KOREA, REPUBLIC OF
       Yang, Eun Young, Suwon-city, KOREA, REPUBLIC OF
       Jeon, Eun Kyung, Yongin-city, KOREA, REPUBLIC OF
       Kim, Chang Ju, Suwon-city, KOREA, REPUBLIC OF
PA
       CJ CORPORATION, Seoul, KOREA, REPUBLIC OF, 100-749 (non-U.S.
       corporation)
       US 20070122480
                            A1 20070531
A1 20040925 (10)
       US 2004-574337
AT
       WO 2004-KR2496
                                 20040925
                                20060509 PCT 371 date
PRAI
       KR 2003-67588
                            20030929
       KR 2004-77158
                            20040924
DT
       Utility
FS
       APPLICATION
LN.CNT 799
INCL
       INCLM: 424/472.000
NCL
       NCLM: 424/472.000
              A61K0009-24 [I,A]
IC
       IPCI
       IPCR
              A61K0009-24 [I,C]; A61K0009-24 [I,A]; A61K0009-22 [I,C*];
              A61K0009-22 [I,A]
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
-> d his
     (FILE 'HOME' ENTERED AT 22:22:24 ON 13 MAR 2009)
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           1288 S TAMSULOSIN
            226 S TAMSULOSIN/CLM
        1224605 S POLYMER?
L.4
         432182 S POLYMER?/CLM
        1287207 S (CALCIUM OR POTASSIUM OR SODIUM OR MAGNESIUM)
L5
L6
         305082 S (CALCIUM OR POTASSIUM OR SODIUM OR MAGNESIUM)/CLM
L7
       2063906 S (MULTILAYER OR LAYER)
1.8
        810896 S (MULTILAYER OR LAYER)/CLM
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L10
         27197 S CAPSULE?/CLM
L11
            242 S L1 AND L4
            897 S L1 AND L3
L13
            888 S L5 AND L12
600 S L7 AND L13
L14
            493 S L9 AND L14
L15
L16
             50 S L2 AND L4
             33 S L6 AND L16
L18
              6 S L8 AND L17
L19
              2 $ L10 AND L18
=> d 115 483-493
L15 ANSWER 483 OF 493 USPAT2 on STN
    Text
AN
       2003:100119 USPAT2
TI
       Ouinazolone derivatives as alpha 1A/B adrenergic receptor antagonists
       Becker, Cyrus Kephra, Menlo Park, CA, UNITED STATES
       Caroon, Joan Marie, Mountain View, CA, UNITED STATES
       Melville, Chris Richard, Palo Alto, CA, UNITED STATES
Padilla, Fernando, Fremont, CA, UNITED STATES
       Pfister, Jurg Roland, Los Altos, CA, UNITED STATES
       Zhang, Xiaoming, Campbell, CA, UNITED STATES
PA
       Syntex (U.S.A.) LLC, Palo Alto, CA, UNITED STATES (U.S. corporation)
PΙ
       US 6900220
                            B2 20050531
       US 2002-40319
                                20020102 (10)
ΑI
PRAT
       US 2001-259337P
US 2001-325267P
                            20010102 (60)
                            20010927 (60)
       Utility
DT
FS
       GRANTED
LN.CNT 2798
INCL
       INCLM: 514/266.210
       INCLS: 514/234.200; 514/234.500; 514/249.000; 514/252.170; 514/264.100;
              514/266.200; 544/116.000; 544/117.000; 544/279.000; 544/284.000;
               544/350.000
NCL
       NCLM:
              514/266.210; 514/223.200
              514/234.200; 514/234.500; 514/249.000; 514/252.170; 514/264.100;
       NCLS:
              514/266.200; 544/116.000; 544/117.000; 544/279.000; 544/284.000;
              544/350.000: 544/012.000
       İCM
              A61K031-517
              A61K031-535; C07D487-00; C07D417-00; C07D471-00
       ICS
              C07D0285-22 [ICM, 7]; C07D0285-00 [ICM, 7, C*]; A61K0031-549
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               [ICS.7]; A61K0031-517 [ICS,7]
       IPCI-2 A61K0031-517 [ICM, 7]; A61K0031-535 [ICS, 7]; C07D0487-00 [ICS, 7];
              C07D0417-00 [ICS,7]; C07D0471-00 [ICS,7]
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              A61K0031-519 [I,A]; A61K0031-549 [I,C*]; A61K0031-549 [I,A];
              C07D0401-00 [I,C*]; C07D0401-04 [I,A]; C07D0401-14 [I,A];
              C07D0403-00 [I,C*]; C07D0403-14 [I,A]; C07D0405-00 [I,C*];
              C07D0405-14 [I,A]; C07D0417-00 [I,C*]; C07D0417-04 [I,A];
              C07D0417-14 [I,A]; C07D0471-00 [I,C*]; C07D0471-04 [I,A];
              C07D0471-14 [I,A]; C07D0487-00 [I,C*]; C07D0487-04 [I,A];
              C07D0491-00 [I,C*]; C07D0491-04 [I,A]
       514/234.2; 514/234.5; 514/249; 514/252.17; 514/264.1; 514/266.2;
       514/266.21; 544/116; 544/117; 544/279; 544/284; 544/350
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L15 ANSWER 484 OF 493 USPAT2 on STN
Full Text
AN
       2003:92739 USPAT2
       Solid carriers for improved delivery of hydrophobic active ingredients
       in pharmaceutical compositions
       Patel, Mahesh V., Salt Lake City, UT, United States
ΤN
       Chen, Feng-Jing, Salt Lake City, UT, United States
Lipocine, Inc., Salt Lake City, UT, United States (U.S. corporation)
PA
       US 6569463
                           B2 20030527
       US 2001-800593
AΤ
                                 20010306 (9)
RLI
       Division of Ser. No. US 1999-447690, filed on 23 Nov 1999, now patented,
       Pat. No. US 6248363
       Utility
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GRANTED
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TMCI.
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        INCLS: 424/422.000; 424/427.000; 424/430.000; 424/433.000; 424/434.000;
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                424/463.000; 424/464.000; 424/465.000; 424/466.000; 424/470.000;
424/474.000; 424/476.000; 424/482.000; 424/490.000; 424/489.000;
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                514/779.000
               424/497.000; 424/465.000
NCL
        NCLM:
        NCLS: 424/422.000; 424/427.000; 424/430.000; 424/433.000; 424/434.000;
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                424/463.000; 424/464.000; 424/465.000; 424/466.000; 424/470.000; 424/474.000; 424/474.000; 424/474.000; 424/489.000; 424/489.000; 424/490.000;
                424/498.000: 514/773.000: 514/779.000: 514/784.000: 514/785.000:
                514/786.000; 977/906.000; 977/927.000
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        ICS
                A61K009-28; A61K009-32; A61K009-52; A61K009-56; A61K009-58
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                A61K0009-14 [I,A]; A61K0009-16 [I,C*]; A61K0009-16 [I,A];
                A61K0009-20 [I,C*]; A61K0009-20 [I,A]; A61K0009-30 [I,C*];
                A61K0009-32 [I,A]; A61K0009-48 [I,C*]; A61K0009-48 [I,A];
                A61K0009-50 [I,C*]; A61K0009-50 [I,A]; A61K0009-51 [I,C*]; A61K0009-51 [I,A]; A61K0009-52 [I,A];
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                A61K0031-403 [I,C*]; A61K0031-404 [I,A]; A61K0031-415 [I,C*];
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                A61K0031-4427 [I,C*]; A61K0031-4439 [I,A]; A61K0031-472 [I,C*];
                A61K0031-4725 [I,A]; A61K0031-519 [I,C*]; A61K0031-522 [I,A];
                A61K0031-57 [I,C*]; A61K0031-57 [I,A]; A61K0031-64 [I,C*];
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                A61K0038-23 [I,C*]; A61K0038-23 [I,A]; A61K0047-02 [I,C*];
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                A61K0047-14 [I,C*]; A61K0047-14 [I,A]; A61K0047-22 [I,C*];
                A61K0047-22 [I,A]; A61K0047-26 [I,C*]; A61K0047-26 [I,A];
                A61K0047-28 [I,C*]; A61K0047-28 [I,A]; A61K0047-32 [I,C*];
                A61K0047-32 [I,A]; A61K0047-36 [I,C*]; A61K0047-36 [I,A];
                A61K0047-38 [I,C*]; A61K0047-38 [I,A]; A61K0047-44 [I,C*];
                A61K0047-44 [I,A]; A61P0001-00 [I,C*]; A61P0001-04 [I,A];
                A61P0003-00 [I,C*]; A61P0003-04 [I,A]; A61P0003-06 [I,A];
                A61P0003-10 [I,A]; A61P0005-00 [I,C*]; A61P0005-16 [I,A];
                A61P0005-24 [I,A]; A61P0005-40 [I,A]; A61P0007-00 [I,C*];
                A61P0007-02 [I,A]; A61P0007-10 [I,A]; A61P0009-00 [I,C*];
                A61P0009-04 [I,A]; A61P0009-06 [I,A]; A61P0009-10 [I,A];
                A61P0009-12 [I,A]; A61P0013-00 [I,C*]; A61P0013-08 [I,A];
                A61P0015-00 [I,C*]; A61P0015-10 [I,A]; A61P0017-00 [I,C*];
                A61P0017-12 [I,A]; A61P0019-00 [I,C*]; A61P0019-06 [I,A];
                A61P0019-10 [I,A]; A61P0021-00 [I,C*]; A61P0021-02 [I,A]; A61P0025-00 [I,C*]; A61P0025-04 [I,A]; A61P0025-06 [I,A];
                A61P0025-08 [I,A]; A61P0025-16 [I,A]; A61P0025-20 [I,A];
                A61P0025-22 [I,A]; A61P0025-26 [I,A]; A61P0025-28 [I,A];
                A61P0029-00 [I,C*]; A61P0029-00 [I,A]; A61P0031-00 [I,C*];
                A61P0031-04 [I,A]; A61P0031-10 [I,A]; A61P0031-12 [I,A];
                A61P0033-00 [I,C*]; A61P0033-06 [I,A]; A61P0033-10 [I,A];
        A61F0035-00 [1,C]; A61F0035-00 [1,A]; A61F0037-00 [1,C];
A61F0035-00 [1,C]; A61F0035-00 [1,A]; A61F0037-00 [1,C];
A61F0037-06 [1,A]; A61F0047-00 [1,C]; A61F0043-00 [1,A]
424/422; 424/433; 424/436; 424/435; 424/40; 424/451; 424/452; 424/464;
EXF
        424/465; 424/482; 424/489; 424/490; 424/480; 424/463; 424/470; 424/497;
        424/498; 424/476; 424/427; 424/430; 424/434; 424/441; 424/466; 424/474
```

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

```
AN
       2003:78121 USPAT2
ΤI
       Modified release formulations containing a hypnotic agent
       Lemmens, Jacobus M., Mook, NETHERLANDS
       van den Heuvel, Dennie J. M., Boxmeer, NETHERLANDS
       Platteeuw, Johannes J., s'Hertogenbosch, NETHERLANDS
       van Dalen, Frans, Nijmegen, NETHERLANDS
Synthon BV, Nijmegen, NETHERLANDS (non-U.S. corporation)
PA
                             B2 20031028
PΤ
       US 6638535
AΙ
       US 2001-833662
                                 20010413 (9)
       US 2000-196939P
                             20000413 (60)
PRAI
       Utility
DT
       GRANTED
LN.CNT 902
INCL
       INCLM: 424/489.000
       INCLS: 424/490.000; 424/464.000; 424/465.000; 514/300.000; 514/781.000
NCL
       NCLM: 424/489.000
       NCLS: 424/464.000; 424/465.000; 424/490.000; 514/300.000; 514/781.000
       ICM
               A61K009-20
               A61K009-14; A61K009-16; A61K031-44; A61K047-00
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               A61K0009-14 [ICM, 7]
       IPCI-2 A61K0009-20 [ICM, 7]; A61K0009-14 [ICS, 7]; A61K0009-16 [ICS, 7];
               A61K0031-44 [ICS, 7]; A61K0047-00 [ICS, 7]
               A61K0009-16 [I,C*]; A61K0009-16 [I,A]; A61K0009-26 [I,C*];
       IPCR
               A61K0009-26 [I,A]; A61K0031-4353 [I,C*]; A61K0031-437 [I,A];
               A61K0031-519 [I,C*]; A61K0031-519 [I,A]
EXF 424/489; 424/490; 424/464; 424/465; 514/300; 514/781 CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L15 ANSWER 486 OF 493 USPAT2 on STN
Full Text
AN
       2002:308400 USPAT2
TI
       Heterocyclic compounds their production and use
IN
       Tasaka, Akihiro, Suita, JAPAN Hitaka, Takenori, Takarazuka, JAPAN
       Matsutani, Etsuva, Suita, JAPAN
       Takeda Chemical Industries, Ltd., Osaka, JAPAN (non-U.S. corporation)
PA
PT
       IIS 6716863
                             B2 20040406
       WO 2001077107
                                  20011018
       US 2001-889974
ΑI
                                  20010724 (9)
       WO 2001-JP2937
                                  20010405
                             20000407
PRAI
       JP 2000-106836
DT
       Utility
       GRANTED
LN.CNT 3144
INCL
       INCLM: 514/374.000
       INCLS: 548/235.000
       NCLM: 514/374.000
NCL
       NCLS: 548/235.000
       ICM
               A61K031-422
       ICS
               C07D413-12
               C07D0413-02 [ICM, 7]; C07D0413-00 [ICM, 7, C*]; A61K0031-422 [ICS, 7]
       IPCI-2 A61K0031-422 [ICM,7]; C07D0413-12 [ICS,7]; C07D0413-00 [ICS,7,C*]
               A61P0035-00 [I,C*]; A61P0035-00 [I,A]; C07D0263-00 [I,C*];
               C07D0263-32 [I,A]; C07D0413-00 [I,C*]; C07D0413-12 [I,A]
EXF 514/374; 548/235
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L15 ANSWER 487 OF 493 USPAT2 on STN
Full Text
AN
       2002:301622 USPAT2
TT
       Benzimidazoles that are useful in treating male sexual dysfunction
       Cowart, Marlon D., Round Lake Beach, IL, UNITED STATES
       Bhatia, Pramila A., Libertyville, IL, UNITED STATES
       Daanen, Jerome F., Racine, WI, UNITED STATES
Stewart, Andrew O., Libertyville, IL, UNITED STATES
       Patel, Meena V., Green Oaks, IL, UNITED STATES
       Kolasa, Teodozyj, Lake Villa, IL, UNITED STATES
       Brioni, Jorge D., Vernon Hills, IL, UNITED STATES
       Rohde, Jeffrey, Evanston, IL, UNITED STATES
```

```
PΑ
       Abbott Laboratories, Abbott Park, IL, UNITED STATES (U.S. corporation)
       US 7022728
                           B2 20060404
AΙ
       US 2002-94265
                                20020308 (10)
PRAI
       US 2001-340452P
                           20011214 (60)
       US 2001-296078P
                           20010605 (60) 20010309 (60)
       US 2001-274805P
       Utility
FS
       GRANTED
LN.CNT 2990
INCL
       INCLM: 514/395.000
       INCLS: 514/394.000; 514/393.000; 514/252.140; 514/254.060; 514/253.010
NCT.
       NCLM: 514/395.000; 514/252.190
              514/252.140; 514/253.010; 514/254.060; 514/393.000; 514/394.000; 514/253.090; 514/254.030
A61K0031-496 [ICM,7]
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       IPCI-2 A61K0031-415 [I,A]; A61K0031-495 [I,A]; A61K0031-50 [I,A]
       IPCR
              A61K0031-496 [I,C*]; A61K0031-496 [I,A]; A61K0031-415 [I,A];
              A61K0031-415 [I,C]; A61K0031-495 [I,C]; A61K0031-495 [I,A];
              A61K0031-50 [I,C]; A61K0031-50 [I,A]
       514/255; 514/258; 514/394; 514/393; 514/359; 514/362; 514/363; 514/385;
EXE
       514/395; 514/252.14; 514/254.06; 514/253.01
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L15 ANSWER 488 OF 493 USPAT2 on STN
AN
       2002:288146 USPAT2
TI
       As-needed administration of tricyclic and other non-SRI antidepressant
       drugs to treat premature ejaculation
IN
       Tam, Peter, Redwood City, CA, UNITED STATES
       Gesundheit, Neil, Los Altos, CA, UNITED STATES
       Wilson, Leland F., Menlo Park, CA, UNITED STATES
       Vivus, Inc., Mountain View, CA, UNITED STATES (U.S. corporation)
PA
ΡI
       US 6946141
                           B2 20050920
       US 2001-996407
ΑI
                                20011121 (9)
RLI
       Continuation-in-part of Ser. No. US 2000-721412, filed on 21 Nov 2000,
       Pat. No. US 6495154
DT
       Utility
FS
       GRANTED
LN.CNT 1495
       INCLM: 424/423.000
INCL
       INCLS: 424/434.000; 424/435.000; 424/443.000; 424/449.000; 424/451.000;
              424/464.000; 424/045.000; 424/046.000
NCL
       NCLM:
              424/423.000; 514/278.000
              424/045.000; 424/046.000; 424/434.000; 424/435.000; 424/443.000;
       NCLS:
              424/449.000; 424/451.000; 424/464.000
       [7]
       ICM
              A61F002-02
       ICS
              A61F013-02; A61K009-48; A61K009-70; A61K009-04
       IPCI
              A61K0031-44 [ICM, 7]
       IPCI-2 A61F0002-02 [ICM, 7]; A61F0013-02 [ICS, 7]; A61K0009-48 [ICS, 7];
              A61K0009-70 [ICS, 7]; A61K0009-04 [ICS, 7]
              A61K0009-02 [I,C*]; A61K0009-02 [I,A]; A61K0009-08 [I,C*];
       IPCR
              A61K0009-08 [I,A]; A61K0009-12 [I,C*]; A61K0009-12 [I,A];
              A61K0009-19 [I,C*]; A61K0009-19 [I,A]; A61K0009-20 [I,C*];
              A61K0009-20 [I,A]; A61K0009-46 [I,C*]; A61K0009-46 [I,A];
              A61K0031-135 [I,C*]; A61K0031-135 [I,A]; A61K0031-136 [I,C*];
              A61K0031-136 [I,A]; A61K0031-403 [I,C*]; A61K0031-404 [I,A];
              A61K0031-55 [I,C*]; A61K0031-55 [I,A]; A61K0031-551 [I,C*]; A61K0031-551 [I,A]; A61K0031-553 [I,A]; A61K0031-553 [I,A];
              A61K0031-554 [I,C*]; A61K0031-554 [I,A]; A61K0045-00 [I,C*];
              A61K0045-00 [I,A]; A61K0045-06 [I,A]; A61K0047-34 [I,C*];
              A61K0047-34 [I,A]; A61P0015-00 [I,C*]; A61P0015-00 [I,A];
              A61P0043-00 [I,C*]; A61P0043-00 [I,A]
EYE
       424/423; 424/434; 424/435; 424/443; 424/449; 424/451; 424/464; 424/45;
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L15 ANSWER 489 OF 493 USPAT2 on STN
AN
       2002:67175 USPAT2
```

Administration of phosphodiesterase inhibitors for the treatment of premature ejaculation $% \left\{ 1,2,\ldots ,2,3,\ldots ,2,1$

```
ΤN
            Wilson, Leland F., Menlo Park, CA, United States
            Doherty, Jr., Paul C., Cupertino, CA, United States
            Place, Virgil A., Kawaihae, HI, United States
            Smith, William L., Montclair, NJ, United States
            Abdel-Hamid Abdou Ali, Ibrahim AbouBakr, Mansoura, EGYPT
Vivus, Inc., Mountain View, CA, United States (U.S. corporation)
PA
                                                B2 20020611
PΙ
            US 6403597
ÀΙ
            US 2001-888250
                                                        20010621 (9)
RLT
            Continuation-in-part of Ser. No. US 1999-467094, filed on 10 Dec 1999
            Continuation-in-part of Ser. No. US 1998-181070, filed on 27 Oct 1998,
            now patented, Pat. No. US 6037346, issued on 14 Mar 2000
            Continuation-in-part of Ser. No. US 1997-958816, filed on 28 Oct 1997,
            now abandoned
            Utility
            GRANTED
FS
LN.CNT 2030
INCL
            INCLM: 514/256.000
            NCLM: 514/256.000; 514/001.000
NCL
IC
            [7]
            ICM
                         A61K031-50
                         A61K0031-00 [ICM, 7]
            IPCI-2 A61K0031-50 [ICM, 7]
            IPCR
                         A61K0009-00 [I,C*]; A61K0009-00 [I,A]; A61K0031-00 [I,C*];
                         A61K0031-00 [I,A]; A61K0031-343 [I,C*]; A61K0031-343 [I,A];
                         A61K0031-381 [I,C*]; A61K0031-381 [I,A]; A61K0031-40 [I,C*];
                         A61K0031-40 [I,A]; A61K0031-4015 [I,C*]; A61K0031-4015 [I,A];
                         A61K0031-4164 [I,C*]; A61K0031-4164 [I,A]; A61K0031-4166 [I,A];
                         AGIRO031-4104 [L/C], AGIRO031-4204 [L/A], AGIRO031-4305 [L/C*];
AGIRO031-437 [L/A]; AGIRO031-4427 [L/C*]; AGIRO031-4439 [L/A];
AGIRO031-444 [L/A]; AGIRO031-4704 [L/C*];
AGIRO031-4709 [L/C*];
AGIRO03
                         A61K0031-4745 [I,A]; A61K0031-496 [I,C*]; A61K0031-496 [I,A];
                         A61K0031-50 [I,C*]; A61K0031-50 [I,A]; A61K0031-502 [I,C*]; A61K0031-502 [I,A]; A61K0031-5025 [I,A];
                         A61K0031-505 [I,C*]; A61K0031-505 [I,A]; A61K0031-519 [I,C*];
                         A61K0031-519 [I,A]; A61K0031-52 [I,A]; A61K0031-522 [I,A]; A61K0031-5575 [I,C*]; A61K0031-558 [I,A]; A61K0031-549 [I,C*]; A61K0031-554 [I,A]; A61K0031-554 [I,A]; A61K0031-554 [I,A];
                         A61K0045-00 [I,C*]; A61K0045-06 [I,A]
            514/258
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L15 ANSWER 490 OF 493 USPAT2 on STN
        Text
            2002:55008 USPAT2
AN
            Clear oil-containing pharmaceutical compositions containing a
TI
            therapeutic agent
            Chen, Feng-Jing, Salt Lake City, UT, United States
IN
            Patel, Mahesh V., Salt Lake City, UT, United States
            Fikstad, David T., Salt Lake City, UT, United States
            Lipocine, Inc., Salt Lake City, UT, United States (U.S. corporation)
PA
ΡI
            US 6761903
                                                 B2 20040713
            US 2001-877541
                                                         20010608 (9)
ΑI
RLI
            Continuation-in-part of Ser. No. US 1999-345615, filed on 30 Jun 1999,
            now patented, Pat. No. US 6267985 Continuation-in-part of Ser. No. US
            2000-751968, filed on 29 Dec 2000, now patented, Pat. No. US 6458383
            Continuation-in-part of Ser. No. US 1999-375636, filed on 17 Aug 1999,
            now patented, Pat. No. US 6309663
DT
            Utility
            GRANTED
FS
LN.CNT 3614
INCL
            INCLM: 424/451.000
            INCLS: 424/043.000; 424/433.000; 424/436.000; 424/441.000; 424/445.000;
                         424/455.000; 424/456.000; 424/458.000; 424/463.000; 424/464.000;
                          424/465.000; 424/489.000; 424/490.000; 424/725.000; 514/772.200;
                        514/772.300; 514/777.000; 514/779.000; 514/781.000; 514/783.000; 514/784.000; 514/785.000; 514/786.000; 514/937.000; 514/944.000 424/451.000; 514/054.000
NCL
            NCLM:
                        424/043.000; 424/433.000; 424/436.000; 424/441.000; 424/445.000;
            NCLS:
                         424/455.000; 424/456.000; 424/458.000; 424/463.000; 424/464.000;
                         424/465.000; 424/489.000; 424/490.000; 424/725.000; 514/772.200;
                         514/772.300; 514/777.000; 514/779.000; 514/781.000; 514/783.000;
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514/784.000; 514/785.000; 514/786.000; 514/937.000; 514/944.000;
               424/727.000; 424/731.000; 424/750.000; 424/757.000
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       TCM
              A61K009-08
       ICS
               A61K009-10; A61K009-14; A61K009-20; A61K009-48
               A61K0031-715 [ICM, 7]; A61K0035-78 [ICS, 7]
       IPCI
       IPCI-2 A61K0009-08 [ICM, 7]; A61K0009-10 [ICS, 7]; A61K0009-14 [ICS, 7];
               A61K0009-20 [ICS, 7]; A61K0009-48 [ICS, 7]
              A61K0009-48 [I,C*]; A61K0009-48 [I,A]; A61K0036-185 [I,C*];
              A61K0036-47 [I,A]
       424/451; 424/450; 424/433; 424/436; 424/441; 424/443; 424/445
EXE
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L15 ANSWER 491 OF 493 USPAT2 on STN
Full Text
AN
       2002:27516 USPAT2
ΤI
       Compounds and methods to increase plasma HDL cholesterol levels and
       improve HDL functionality
       Luchoomun, Jayraz, Lilburn, GA, UNITED STATES
       Meng, Charles Q., Alpharetta, GA, UNITED STATES
       Saxena, Uday, Atlanta, GA, UNITED STATES
       Sikorski, James A., Alpharetta, GA, UNITED STATES
PA
       Atherogenics, Inc., Alpharetta, GA, UNITED STATES (U.S. corporation)
PΙ
       US 6881860
                           B2 20050419
       US 2001-833407
ΑI
                                20010411 (9)
PRAI
       US 2000-196201P
                            20000411 (60)
DT
       Utility
FS
       GRANTED
LN.CNT 3107
       INCLM: 562/426.000
INCL
       INCLS: 514/568.000
       NCLM: 562/426.000; 514/517.000
NCL
       NCLS: 514/571.000; 514/649.000; 558/037.000; 564/347.000
       [7]
       İCM
              C07C321-00
       IPCI
               A61K0031-255 [ICM, 7]; A61K0031-21 [ICM, 7, C*]; A61K0031-192
               [ICS, 7]; A61K0031-185 [ICS, 7, C*]; A61K0031-145 [ICS, 7]
       IPCI-2 C07C0321-00 [ICM, 71
       TPCR
              C07C0323-00 [I,C*]; C07C0323-20 [I,A]; G01N0033-92 [I,C*];
               G01N0033-92 [I,A]
       562/426; 514/568
EXE
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L15 ANSWER 492 OF 493 USPAT2 on STN
Full Text
AN
       2002:21845 USPAT2
ΤI
       Compositions and methods for improved delivery of hydrophobic agents
IN
       Patel, Mahesh V., Salt Lake City, UT, United States
       Chen, Feng-Jing, Salt Lake City, UT, United States
Lipocine, Inc., Salt Lake City, UT, United States (U.S. corporation)
PA
ΡI
       US 6451339
                           B2 20020917
ΑI
       US 2001-898553
                                20010702 (9)
       Continuation of Ser. No. US 1999-258654, filed on 26 Feb 1999, now
RLI
       patented, Pat. No. US 6294192
DT
       Ütilitv
FS
       GRANTED
LN.CNT 2907
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       INCLM: 424/451.000
       INCLS: 424/450.000: 424/455.000: 424/456.000: 424/463.000: 424/489.000:
               424/499.000; 424/502.000; 424/435.000; 424/464.000; 424/937.000;
               424/938.000; 424/939.000; 514/940.000; 514/941.000; 514/942.000;
               514/943.000; 514/975.000
       NCLM:
NCL.
              424/451.000; 424/400.000
       NCLS:
              424/435.000; 424/450.000; 424/455.000; 424/456.000; 424/463.000;
               424/464.000; 424/489.000; 424/499.000; 424/502.000; 514/937.000;
               514/938.000; 514/939.000; 514/940.000; 514/941.000; 514/942.000; 514/943.000; 514/975.000
TC:
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       ICM
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       IPCI
              A61K0009-00 [ICM, 7]
       IPCI-2 A61K0009-127 [ICM, 7]
       IPCR A61K0009-48 [I,C*]; A61K0009-48 [I,A]; A61K0031-57 [I,C*];
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A61K0031-57 [I,A]; A61K0038-12 [I,C*]; A61K0038-13 [I,A]
EXF
       424/450; 424/451; 424/455; 424/456; 424/463; 424/489; 424/499; 424/502;
       424/435; 424/464; 514/937; 514/938; 514/939; 514/940; 514/941; 514/943;
       514/975
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L15 ANSWER 493 OF 493 USPAT2 on STN
     Text
AN
       2001:229703 USPAT2
       Co-administration of a prostaglandin and an androgenic agent in the
       treatment of female sexual dysfunction
TM
       Place, Virgil A., Kawaihae, HI, United States
       Wilson, Leland F., Menlo Park, CA, United States
Doherty, Jr., Paul C., Cupertino, CA, United States
       Hanamoto, Mark S., Belmont, CA, United States
       Spivack, Alfred P., Menlo Park, CA, United States
       Gesundheit, Neil, Los Altos, CA, United States
       Bennett, Sean R., Denver, CO, United States
PA
       Vivus, Inc., Mountain View, CA, United States (U.S. corporation)
ΡI
       US 6593313
                             B2 20030715
ΑI
       US 2001-905458
                                  20010713 (9)
RLI
       Continuation of Ser. No. US 2000-539484, filed on 30 Mar 2000, now
       patented, Pat. No. US 6306841 Continuation of Ser. No. US 1998-181316,
       filed on 27 Oct 1998, now abandoned Continuation-in-part of Ser. No. US
       1997-959064, filed on 28 Oct 1997, now patented, Pat. No. US 5877216
       Continuation-in-part of Ser. No. US 1997-959057, filed on 28 Oct 1997,
       now abandoned
       Utility
FS
       GRANTED
LN.CNT 1331
INCL
       INCLM: 514/108.000
       NCLM: 514/108.000; 514/530.000
NCL
       NCLS: 514/288.000; 514/573.000
       [7]
       ICM
               A61K031-19
       ICS
               A61K031-557
       IPCI
               A61K0031-5575 [ICM, 7]; A61K0031-557 [ICM, 7, C*]; A61K0031-48
       IPCI-2 A61K0031-19 [ICM, 7]; A61K0031-185 [ICM, 7, C*]; A61K0031-557
               [ICS, 7]
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               A61K0009-00 [I,C*]; A61K0009-00 [I,A]; A61K0009-02 [N,C*];
               A61K0009-02 [N,A]; A61K0031-00 [I,C*]; A61K0031-00 [I,A];
               A61K0031-15 [I,C*]; A61K0031-15 [I,A]; A61K0031-21 [I,C*]; A61K0031-21 [I,A]; A61K0031-21 [I,A]; A61K0031-28 [I,C*]; A61K0031-29 [I,A]; A61K0031-29 [I,A]; A61K0031-28 [I,C*];
               A61K0031-5377 [I,A]; A61K0031-557 [I,C*]; A61K0031-557 [I,A]; A61K0031-5575 [I,A]; A61K0031-5585 [I,A]; A61K0031-56 [I,C*];
               A61K0031-56 [I,A]; A61K0045-00 [I,C*]; A61K0045-06 [I,A]
EXF
       514/573
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
=> d 115 kwic 492
L15 ANSWER 492 OF 493 USPAT2 on STN
       The POE-POP block copolymers are a unique class of polymeric
DETD
       surfactants. The unique structure of the surfactants, with hydrophilic
       POE and hydrophobic POP moieties in well-defined ratios and positions,
       provides. . . Synperonic PE series (ICI); Pluronic® series (BASF), Emkalyx, Lutrol (BASF), Supronic, Monolan, Pluracare, and
       Plurodac. The generic term for these polymers is "poloxamer" (CAS
       9003-11-6). These polymers have the formula:
DETD
              . use in the present invention. Preferred anionic surfactants
       include fatty acid salts and bile salts. Specifically, preferred ionic
       surfactants include sodium oleate, sodium lauryl sulfate, sodium
       lauryl sarcosinate, sodium dioctyl sulfosuccinate, sodium cholate,
       and sodium taurocholate. Examples of such surfactants are shown in
       Table 18 below. For simplicity, typical counterions are shown in the
       entries. . . in the art, however, that any bioacceptable counterion
       may be used. For example, although the fatty acids are shown as sodium
       salts, other cation counterions can also be used, such as alkali metal
       cations or ammonium. Unlike typical non-ionic surfactants, these. .
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DETD
TABLE 18
Ionic Surfactants
COMPOUND HLB
FATTY ACID SALTS >10
Sodium caproate
Sodium caprylate
Sodium caprate
Sodium laurate
Sodium myristate
Sodium myristolate
Sodium palmitate
Sodium palmitoleate
Sodium oleate 18
Sodium ricinoleate
Sodium linoleate
Sodium linolenate
Sodium stearate
Sodium lauryl sulfate (dodecyl) 40
Sodium tetradecyl sulfate
Sodium lauryl sarcosinate
Sodium dioctyl sulfosuccinate [sodium docusate (Cytec)]
BILE SALTS >10
Sodium cholate
Sodium taurocholate
Sodium glycocholate
Sodium deoxycholate
Sodium taurodeoxycholate
Sodium glycodeoxycholate
Sodium ursodeoxycholate
Sodium chenodeoxycholate
Sodium taurochenodeoxycholate
Sodium glyco cheno deoxycholate
Sodium cholylsarcosinate
Sodium N-methyl taurocholate
PHOSPHOLIPIDS
Egg/Soy lecithin [Epikuron .TM. (Lucas Meyer), Ovothin .TM. (Lucas
Meyer)]
Lyso egg/soy lecithin
Hydroxylated lecithin
Lysophosphatidylcholine
Cardiolipin
Sphingomyelin
Phosphatidylcholine
Phosphatidyl ethanolamine
Phosphatidic acid
Phosphatidyl glycerol
Phosphatidyl serine
PHOSPHORIC ACID ESTERS
Diethanolammonium polyoxyethylene-10. . . fatty alcohol ethoxylates
with phosphoric acid or anhydride
CARBOXYLATES
Ether carboxylates (by oxidation of terminal OH group of fatty
alcohol ethoxylates)
Succinylated monoglycerides [LAMEGIN ZE (Henkel)]
Sodium stearyl fumarate
Stearoyl propylene glycol hydrogen succinate
Mono/diacetylated tartaric acid esters of mono- and diglycerides
Citric acid esters of mono-, diglycerides
Glyceryl-lacto esters of fatty acids (CFR ref. 172.852)
Acyl lactylates:
 lactylic esters of fatty acids
 calcium/sodium stearoyl-2-lactylate
 calcium/sodium stearoyl lactylate
Alginate salts
Propylene glycol alginate
SULFATES AND SULFONATES
Ethoxylated alkyl sulfates
Alkyl benzene sulfones
α-olefin sulfonates
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Acyl isethionates Acyl taurates

Alkyl glyceryl ether sulfonates

Octvl sulfosuccinate disodium

Disodium undecylenamideo-MEA-sulfosuccinate

CATIONIC. .

DETD A third method of determining optical clarity and carrier diffusivity through the aqueous boundary layer is to quantitatively measure the size of the particles of which the dispersion is composed. These measurements can be performed.

DETD analgesics and anti-inflammatory agents, such as aloxiprin, auranofin, azapropazone, benorylate, capsaicin, celecoxib, diclofenac, diflunisal, etodolac, fenbufen, fenoprofen calcium, flurbiprofen, ibuprofen, indomethacin, ketoprofen, ketorolac, leflunomide, meclofenamic acid, mefenamic acid, nabumetone, naprosen, oxaprozin, oxyphenbutazone, phenylbutazone, piroxicam, refocoxib, sulindac, tetrahydrocannabinol, tramadol.

DEID anti-epileptics, such as beclamide, carbamazepine, clonazepam, ethotoin, felbamate, fosphenytoin **sodium**, lamotrigine, methoin, methsuximide, methylphenobarbitone, oxcarbazepine, paramethadione, phenacemide, phenobarbitone, phenytoin, phensuximide, primidone, sulthiame, tiagabine HCl, topiramate, valproic acid, and vigabatrin;
DEID gatrointestinal agents, such as bisacodyl, cimetidine, cisapride,

DETD gastrointestinal agents, such as bisacodyl, cimetidine, cisapride, diphenoxylate HCl, domperidone, famotidine, lansoprazole, loperamide, mesalazine, nizatidine, omeprazole, ondansetron HCl, rabeprazole sodium, ranitidine HCl and sulphasalazine;

DEID muscle relaxants, such as dantrolene sodium and tizanidine HCl; and others, such as becaplermin, donepezil HCl, L-thryroxine, methoxsalen, verteporfrin, physostigmine, pyridostigmine, raloxifene HCl, sibutramine HCl, sildenafil citrate, tacrine, tamsulosin HCl, and tolterodine.

DEID

. a semi-solid dispersion or a solid dispersion. If desired, the compositions may be encapsulated in a hard or soft gelatin capsule, a starch capsule or an enteric coated capsule. The term "enteric coated capsule" as used herein means a capsule coated with a coating resistant to acid; i.e., an acid resistant enteric coating. Although solubilizers are typically used to enhance. of a hydrophobic therapeutic agent, they may also render the compositions more suitable for encapsulation in hard or soft gelatin capsules. Thus, the use of a solubilizer such as those described above is particularly preferred in capsule dosage forms of the pharmaceutical compositions. If present, these solubilizers should be added in amounts sufficient to impart to the.

DETD . . . is mono-modal and narrow. This reduced and more uniform size enables more efficient drug transport through the intestinal aqueous boundary layer, and through the absorptive brush border membrane. More efficient transport to absorptive sites leads to improved and more consistent absorption.

DEID . . . conditions which limit production of lipase, such as pancreatic lipase secretory diseases; and dependence of lipolysis on stomach pH, endogenous calcium concentration, and presence of co-lipase or other digestion enzymes. The lack of lipolysis dependence further provides transport which does not. . .

transport which does not.

DETD . . the pharmaceutical compositions of the present invention allow for faster transport of the hydrophobic therapeutic agent through the acqueous boundary layer.

DETD . . the hydrophobic therapeutic agent. A third solution was prepared with simulated intestinal fluid, plus an additional aliquot of 20 mM sodium taurocholate (a bile salt); this solution is designated SIFB in Table 29. Finally, a fourth solution was prepared with simulated intestinal fluid, 20 mM sodium taurocholate, and 5 mM lecithin; this solution is designated SIFBL. The 20 mM bile salt and 5 mM lecithin concentrations.

Cyclosporine 0.140 g Cremophor RH-40 0.41 g Arlacel 186 0.29 g Sodium taurocholate 0.26 g Propylene glycol 0.46 g CLM What is claimed is:

^{. .} lecithins; lysolecithin and hydrogenated lysolecithins;

lysophospholipids and derivatives thereof; phospholipids and derivatives thereof; salts of alkylsulfates; salts of fatty acids; **sodium** docusate; and mixtures thereof.

- CLM What is claimed is:
 - and salts, analogues, and derivatives thereof; lecithins, lysolecithin, phospholipids, lysophospholipids and derivatives thereof; salts of alkylsulfates; salts of fatty acids; sodium docusate; acyl lactylates; monoacetylated tartaric acid esters of monoglycerides, monoacetylated tartaric acid esters of diglycerides, diacetylated tartaric acid esters of diglycerides, diacetylated tartaric acid esters of
- CLM What is claimed is: 77. A dosage form comprising a capsule filled with the pharmaceutical formulation of any one of claims 69, 70, 71 or 72.
- CLM What is claimed is:
 78. The dosage form of claim 77, wherein the capsule is a hard gelatin capsule, a soft gelatin capsule, a starch capsule or an enteric coated capsule.
- CLM What is claimed is: 88. The formulation of claim 1, wherein the formulation is contained in a capsule.
- CLM What is claimed is:
 95. The formulation of claim 89, wherein the formulation is contained in a capsule.
- CLM What is claimed is: 105. The formulation of claim 104, wherein the formulation is contained in a capsule.
- CLM What is claimed is: 107. The formulation of claim 106, wherein the formulation is contained in a capsule.
- CLM What is claimed is: 117. The formulation of claim 116, wherein the formulation is contained in a cassule.

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ST IN U.S. DOLLARS
SINCE FILE TOTAL
FULL ESTIMATED COST
42.32
43.86

STN INTERNATIONAL LOGOFF AT 22:34:21 ON 13 MAR 2009